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(54) Title: INK JET RECORDING PAPER INCORPORATING NOVEL PRECIPITATED CALCIUM CARBONATE PIGMENT		
(57) Abstract An improved thermal ink jet recording paper, incorporating heat aged and/or milled precipitated calcium carbonate ("PCC") and a binder, is disclosed. The heat aging and milling processes produce discrete particles of PCC with a surface morphology and chemistry that enhances the ability of the PCC to bind to ink jet ink without binding so strongly that the color of the ink is changed. Ink jet recording papers incorporating the PCC and binders of the present invention have been produced that are superior to multipurpose office papers with reduced feathering, spreading, and penetration or backside showthrough, as well as improved optical density, dry time, and water fastness. The ink jet recording paper of the present invention is comparable to silica based commercial papers, at a fraction of the cost.		

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INK JET RECORDING PAPER INCORPORATING
NOVEL PRECIPITATED CALCIUM CARBONATE PIGMENT

Field of the Invention

The present invention relates to novel heat
5 aged and milled precipitated calcium carbonate pigments
for use in ink jet recording papers. More
particularly, the present invention relates to an ink
jet recording paper which incorporates such pigments
and at least one water soluble binder to impart
10 enhanced print quality. The invention also relates to
methods of applying these components to paper, and to
methods of producing the novel pigments.

Background of the Invention

The thermal ink jet process applies a dilute
15 aqueous ink onto the surface of a paper by heating a
small volume of the ink in a small chamber with an
orifice that is directed at the recording paper. The
small volume of ink that is heated rapidly reaches its
boiling point, and the steam bubble formed propels a
20 tiny drop of liquid ink at the paper, where the drop
produces a single dot in a dot matrix that forms a
character or image on the sheet. This process requires
an ink that is low in solids and high boiling
components so that it is capable of boiling rapidly
25 without leaving a residue that can foul the heating
element, and clog the orifice. Therefore, up to 96
percent of ink jet printer ink is a mixture of water
and low molecular weight glycols. Although such an ink
boils quickly when heated to ensure rapid printing, and

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is not prone to clog, it results in an applied ink that is very mobile and slow to dry. Therefore, good print quality can be obtained only if the ink colorant or dye remains on or near the outer surface of the paper, and
5 does not spread or move from the point at which it was applied.

It is also important that drying occurs rapidly to prevent smearing of the colorant. In printers that are not equipped with heating elements,
10 the water and glycol components of the ink must penetrate into the body of the paper for proper drying of the colorant on the surface. If the colored phase is carried into the paper with the liquid phase as it penetrates into the paper, or if the colorant migrates
15 across the surface of the paper, the quality of the resulting print or image will be poor. Also, dry ink colorant that is not permanently fixed on the paper will blot or run if the printed surface becomes wet or is marked with a highlighter. Therefore, the dry ink
20 should have excellent water and highlighter fastness properties for optimum performance.

In most applications, multipurpose office papers provide inadequate or poor thermal ink jet print quality. This is particularly true where multicolor
25 printing with concomitant superimposed ink applications is utilized. The poor print quality is compounded in printers that apply the colors in one order when the print head moves to the right and the reverse order when the print head moves to the left. Multipurpose
30 office papers often allow the colorant to penetrate into the paper, which results in reduced optical density of the printed image, and increased showthrough on the reverse side of the paper. Multipurpose office papers that are highly sized prevent liquid
35 penetration, leading to higher ink optical density, but, also, excessive feathering and spreading.

One method of improving thermal ink jet print

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quality is to apply a material to the paper surface that binds the ink colorant to the surface, but allows the water/glycol liquid phase to pass into the body of the paper, which speeds drying. However, the ink colorant often is an unsaturated or aromatic organic compound, and if the surface material interacts too strongly with the colorant the color of the ink can change. Therefore, a surface material must be obtained that prevents the ink colorant from penetrating the paper, but does not interact so strongly as to effect the colorant, and cause a color change.

Oshima et al., U.S. Patent No. 4,478,910, discloses a paper base stock that is coated with a high specific surface area colloidal silica pigment and a polyvinyl alcohol binder, where the specific surface area of the colloidal silica is greater than 200 m²/g.

Migamoto, U.S. Patent No. 4,576,867, and Kojima et al., U.S. Patent No. 4,830,911, teach that application of a cationic quaternary ammonium polymer to a paper stock will improve the water fastness of the resulting paper. Application of such a quaternary ammonium polymer in a size press is also disclosed by Malhotra, U.S. Patent No. 5,223,338. However, high quality, commercial thermal ink jet recording paper is produced with off machine coaters, i.e., the paper is not treated in the size press of the paper making machine, but must be removed and coated in an additional step using a separate coating process.

Precipitated calcium carbonate ("PCC") has been disclosed as a useful pigment or filler in ink jet paper, but is typically included in a long list of compounds that also includes natural ground calcium carbonate, a form of the compound that differs greatly from PCC in its physical and chemical properties; e.g., natural ground calcium carbonate particles are far larger than those found in PCC, and are normally anionic, whereas PCC is naturally cationic.

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Kondo et al., U.S. Patent No. 5,320,897, describes coating ink jet recording paper with off-the-shelf pigments and a water soluble binder. The pigments which are disclosed as being useful have an
5 apparent specific gravity in the range of 0.10 to 0.50 g/cm³, and produce a paper with a water contact angle of 45° to 100°.

Tsukisaka et al., U.S. Patent No. 5,007,964, disclose a generic precipitation method incorporating a
10 chelating agent to obtain a porous agglomeration of chain like PCC particles with a BET specific surface area in the range of 25 to 55 m²/g for use in a recording paper. Particles with a BET specific surface area of greater than 55 m²/g, are disclosed as being
15 undesirable because they have a higher density, are not porous, and exhibit a low ability to absorb oil and water. It is further disclosed that PCC can be used as a paste, or as a powder obtained by pulverizing the dried paste, which is disclosed as not affecting the
20 agglomeration of the particles.

Kunesh et al., U.S. Patent No. 5,215,734, disclose an accelerated heat aging process for PCC that produces crystals with a specific surface area of about
3 to about 15 m²/g and an average discrete particle
25 size of about 0.2 to about 0.9 μm. The PCC is used to improve the optical properties of paper.

In view of the above, there is nothing in the prior art to suggest how to improve the print quality of ink jet recording paper. The present invention
30 provides one such solution to this problem.

Summary of the Invention

The present invention relates to an ink jet recording paper with enhanced properties that incorporates novel heat aged and/or milled precipitated
35 calcium carbonate pigments and a binder. The heat aged PCC is prepared by carbonating a slurry of calcium hydroxide to form a slurry of precipitated calcium

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carbonate, screening the slurry to remove impurities, adding an organophosphonate compound, such as ethanolamine bis-(methylenephosphonic acid), to the slurry, and heat aging the PCC slurry in the presence of the organophosphonate to produce an improved pigment for ink jet recording paper.

The ink jet recording paper of the present invention is produced by coating at least one side of a paper base stock with a coating comprising heat aged and/or milled PCC and a binder to form the ink jet recording paper. The heat aged and/or milled precipitated calcium carbonate is made by carbonating a slurry of calcium carbonate, heat aging the PCC in the presence of an organophosphonate compound and/or milling the PCC to produce discrete particles of PCC with a surface morphology and chemistry that enhances the ability of the PCC to bind to the ink jet printer ink colorant.

Detailed Description of the Invention

An ink jet recording paper has been obtained that provides full color ink jet print quality that approaches or surpasses that of the best ink jet specialty paper at a greatly reduced price. The selection of PCC particle size, surface area, surface chemistry, and degree of aggregation, and its combination with an appropriate formulation binder allows each thermal ink jet print characteristic to be individually adjusted and optimized.

The cationic PCCs of the present invention bind ink colorants without the use of expensive cationic resins, and can be manufactured on-site at the paper mill, so that, unlike silica, they do not need to be dried, shipped, and then rewetted for use. The present formulations are designed for application in a metering size press, such as those installed on large, low unit cost paper machines. Therefore, the PCC compositions of the present invention can be applied as

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part of the paper making operation, rather than as a separate, additional step after normal papermaking operations are completed that increases the cost of the process and the finished product. As a metered size
5 press application is typically two sided, both sides of a paper coated on such a press are available for printing, rather than the single sided coating that is found in expensive, high quality commercial papers. Although polyvinyl alcohol is a preferred binder in the
10 present invention, treating an ink jet recording paper with heat aged and/or milled PCC provides a significant improvement in print quality with lower cost binders. Finally, the present formulations provide a paper that is still suitable for photocopying, which allows
15 multiple uses of the paper.

PCC is normally produced by bubbling a gas containing carbon dioxide through an aqueous suspension of calcium hydroxide. Other inorganic materials, in particular, inorganic materials containing aluminum,
20 such as alum, can be coprecipitated with PCC, or can be precipitated onto the surface of the PCC precipitate. It has been unexpectedly discovered, however, that when such a PCC is heat aged and/or milled, treated with an appropriate binder, and then used in a composition for
25 treating thermal ink jet recording paper, a high quality ink jet recording paper is obtained, which provides a print quality that compares favorably with that of high cost, commercial, silica coated thermal ink jet papers, but at a greatly reduced cost.

30 The heat aging and milling processes produce discrete particles of PCC with a surface morphology and chemistry that enhances the ability of the PCC to bind to the ink jet ink colorant without binding so strongly that the color of the ink is changed. Ink jet
35 recording papers incorporating the PCC and binders of the present invention have been produced that are superior to multipurpose office papers with reduced

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feathering, spreading, and penetration or backside showthrough, as well as improved optical density, dry time, and water fastness.

The PCCs of the present invention are produced by the conventional precipitation method of introducing a gas containing carbon dioxide into a suspension of calcium hydroxide. The calcium carbonate precipitate is then heat aged and/or milled before use. The print quality of an ink jet recording paper incorporating such a PCC is controlled by the state of aggregation of the particles and the surface chemistry of the pigment and the binder. The chemical nature of the pigment surface is controlled by the specific synthetic pathway and post synthesis treatment of the pigment.

A PCC particularly useful in the present invention is prepared by a process which comprises introducing carbon dioxide into an aqueous slurry of calcium hydroxide at a temperature of from about 7° to about 18°C. The slurry contains from about 0.02 to 1 weight percent, based on the calcium carbonate equivalent of the calcium hydroxide in the reaction slurry, of an anionic organopolyphosphonate polyelectrolyte (typically, Briquest 221-50A, ethanolamine bis-(methylenephosphonic acid), Albright & Wilson Americas, Inc., Richmond, VA) and from 0 to 10 weight percent of aluminum sulfate octadecahydrate ($\text{Al}_2\text{SO}_4 \cdot 16\text{H}_2\text{O}$). The introduction of carbon dioxide is continued until precipitation of the calcium carbonate is substantially complete. This typically occurs when the conductivity of the carbonated slurry reaches a minimum at a pH of about 10 to 11.

Except for the addition of aluminum sulfate octadecahydrate, the process to this point is essentially that of U.S. Patent No. 4,367,207, the teachings of which are incorporated herein by reference. The addition of aluminum sulfate

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octadecahydrate is disclosed in U.S. Patent No. 4,892,590, the teachings of which are also incorporated herein by reference.

5 Once a PCC of the desired particle size and specific surface area is obtained, the PCC is heat-aged or milled to obtain the PCC characteristics required for a high-quality, low cost ink jet recording paper.

10 Milling breaks pigment aggregates into individual particles, which exposes PCC surfaces that were joined, and are chemically different from the aggregate surface. Milling is typically performed on a PCC with a scalenohedral, acicular, prismatic, or rhombohedral morphology. Heat aging is normally performed on a rhombohedral or prismatic PCC, and not
15 only separates aggregates, but also involves further precipitation on the particle surface. It is believed that the newly precipitated material may be chemically different from the original surface material; e.g., the surface of a heat aged or milled PCC may have a higher
20 calcium-magnesium ratio.

Conventional heat aging, also known as Ostwald ripening, is a process whereby crystals such as calcium carbonate, initially at a higher internal energy state, and having relatively high phase
25 solubilities, undergo a phase transformation by dissolving and redepositing on crystals at a lower internal energy state. The process results in a final crystal product characterized by greater perfection of the crystal lattice structure, a narrower particle size
30 distribution, a greater degree of particle discreteness, and a lower surface energy.

In the present invention, heat aging of PCC starts by ending the reaction of carbon dioxide and calcium hydroxide when the conductivity of the slurry
35 reaches a minimum, which is typically at a pH of about 10-11. The PCC is then screened to remove impurities, and up to about 1 percent by weight of an

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organophosphonate, based on the weight of PCC, is added to control the extent or degree of heat aging. Other chemical agents that are surface active with regard to calcium carbonate will also serve as well as

5 organophosphonates to control the extent or degree of heat aging. Such agents include, but are not limited to the following, sodium polyphosphates, sodium silicates, sodium polyacrylates, various carboxylic acids, such as mono, di, tri, and polycarboxylic acids,

10 and their salts, various polysaccharides, and various gums with repeating carboxylic acid functionalities. The slurry is then heated to a temperature of at least about 75°C. and preferably about 80°C. or higher to heat age the PCC particles. The heat aging step is

15 conducted for up to about 10 hours or more, depending upon the temperature. At temperatures of about 80 to 85°C., the heating is conducted for about 2 to 5 hours. The pH of the slurry rises to approximately 10.5 due to the presence of unreacted calcium hydroxide. The aging

20 reaction is then monitored by measuring the surface area of the PCC at hourly intervals. The heat aged PCC that is recovered as the final product of the process has a rhombohedral or prismatic morphology and a specific surface area of about 60 to about 100 m²/g,

25 preferably about 65 to about 95 m²/g, and most preferably from about 80 to 90 m²/g.

The organophosphonates employed in the present invention are organopolyphosphonates of varying molecular weights commonly used as scale inhibitors,

30 sequesterants, deflocculants, and detergent promoters. Such organophosphonates include, but are not limited to, nitrilo tris (methylene phosphonic acid), ethylenediaminetetra (methylene phosphonic acid), diethylenetriaminepenta (methylene phosphonic acid),

35 hydroxy ethane-1, 1-diphosphonic acid, ethanolamine, ethanolamine bis-(methylenephosphonic acid), N,N-dimethylene phosphonic acid, and hexamethylene-

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diaminetetra (methylene phosphonic acid). The preferred organophosphonate is ethanolamine bis-(methylenephosphonic acid).

The milling of a calcium carbonate may be carried out in either a wet or dry milling process in, for example, a conventional ball mill, jet mill, micro mill, or media mill. The preferred milling is carried out by introducing an aqueous slurry of PCC into a media mill containing glass media of a size from about 0.7 to 0.9 mm. The media mill is equipped with mechanical agitation, and the preferred weight percent solids of the PCC slurry is from about 10 to 30 percent, based on the total weight of the PCC and the water. When media milling is typically performed on PCC with a scalenohedral morphology, a PCC with a specific surface area of from about 10 to about 40 m²/g, preferably from about 15 to about 30 m²/g, and a less elongated crystal shape is produced.

Examples of binders useful for coating compositions for ink jet recording paper are those heretofore conventionally used in the art, and include polyvinyl alcohol ("PVOH") and derivatives thereof, oxidized starch, etherified starch, esterified starch, dextrin and like starches, carboxymethylcellulose, hydroxyethylcellulose and like cellulose derivatives, casein, gelatin, soybean protein, maleic anhydride resin, lattices of usual styrenebutadiene copolymer, methyl methacrylate-butadiene copolymer and like conjugated diene polymers or copolymers, and lattices of acrylate and methacrylate polymers or copolymers and like acrylic polymers, and latex. When required, the coating composition may have further incorporated therein in an amount conventionally used in the art of conventional pigment dispersants, tackifiers, flowability modifiers, defoaming agents, foaming inhibitors, release agents, coloring agents, and the like.

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Ink jet recording papers incorporating the PCC formulations of the present invention have been prepared and tested. The following is a summary of the procedures and testing methods used. Once a PCC is produced, the entire testing process can be categorized into four areas; pigment preparation, formulation with binder, paper coating and processing, and testing. Specific details for each of these procedures are given below.

10 Pigment Preparation:

Pigments to be tested are typically in the form of a slurry or a filter cake. Samples in the form of a slurry are concentrated to the desired solids by vacuum filtration. In some instances, such as with a media milled PCC with a specific surface area of about 10 to 35 m²/g, the slurry solids are not further altered in order to duplicate trial conditions. Once the target formulation solids is set, the pigment is diluted, if necessary, with water and thoroughly mixed. The pigment is characterized by particle size distribution (Sedigraph 5100), specific surface area (Flowsorb), viscosity (Brookfield), and morphology by scanning electron microscopy ("SEM").

The surface area of the product was obtained using a Micromeritics Flowsorb II 2300, which employs BET theory with nitrogen as the absorbing gas. The particle size was determined by a sedimentation technique using a Micromeritics Sedigraph Model 5100 on an aqueous dispersion of the product at about 3 percent and using about 0.1 percent carboxylated polyelectrolyte (Daxad 30) as a dispersant.

30 Formulation:

Typical binders are cationic starch (Filmcote-54[®], National Starch), polyvinyl alcohol or PVOH (Airvol 107[®], Air Products Inc.), and latex (Dow experimental size press latex). When a starch is used as a binder, the dry starch is dispersed in water at

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about 10-25 percent solids, and then cooked in an automated laboratory cooker at about 195°C. for 50-190 minutes. The resulting viscous starch slurry is combined with the pigment, which has been appropriately prepared to attain the target formulation solids, and mixed thoroughly on a Premier mill with a Cowles type open impeller blade. The formulation is mixed for about 5 minutes until a completely homogenous slurry is obtained, and the resulting formulation is characterized by Brookfield viscosity (10, 20, 50, and 100 r.p.m.) and solids.

Polyvinyl alcohol is prepared in a manner similar to that used for starch. The PVOH is hydrated at about 200°C. in a laboratory cooker for 50-190 minutes at 10 percent solids. For the latex binder (50 percent solids), no preparation is necessary before testing. The formulation of these binders with the pigment is the same as with the starch.

Formulation solids for the tests were in the range of about 20-28 percent, with a typical coating formulation having about 20 percent solids.

Paper Coating:

Generally, an unsized base stock with an 81.3-83.0 g/m² basis weight is used in the tests. The paper is cut into 12x17" sheets and secured to the CSD Drawdown Apparatus, which consists of a glass plate (12x17") mounted on metal base with spring clip at the top. A coating formulation is applied with a CSD drawdown rod by placing the rod of choice, which depends on target coat weight, at the top of the paper, adding a thoroughly mixed 10-15 ml sample of the formulation with a 30 ml syringe in a uniform line across the top to the paper, below the rod, and coating the paper by pulling the drawdown rod from top to bottom using light pressure and a constant, steady rate for about 2 seconds.

The coat weight is determine by the stainless

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steel drawdown rods, which are specifically grooved to deliver a predetermined coating volume to the paper surface. Rods with fewer grooves deliver a heavier coat weight, since the spaces between the grooves are wider. In turn, rods with a greater number of more tightly spaced grooves produce lighter coat weights. Typical coat weights are about 2 to 10 g/m².

Once a formulation has been applied, the paper is immediately dried with a hand held heat gun for about 30-60 seconds, and then completely dried at room temperature over a period of about 2-24 hours. The dried papers are then cut into 8 1/2 x 11" sheets for testing.

Paper Testing:

Minimum ink jet print quality specifications have been established by Hewlett Packard Corporation ("HP"). Therefore, most tests utilize HP methods to determine the following print characteristics.

Optical density is a measure of the reflection density of an image. A specific test pattern is printed onto the paper, and the optical densities of pure black, composite black, cyan, magenta, and yellow are measured using a reflection densitometer (Macbeth RD918). The resulting optical densities are compared to minimum HP specifications.

Ink spreading and feathering can both decrease the quality of ink jet print. Ink spreading is defined as the growth or widening of printed areas. Feathering is the wicking of ink, which results in fuzzy images. This is measured by analyzing a specific portion of the same printed pattern used for optical density measurements. The specific portion is evaluated for ink area, spreading, and ink perimeter, feathering. The resulting, digitized pattern is quantitated and compared to a commercial premium ink jet paper. The HP test method for ink spreading and feathering was not used in these tests, since the HP

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test is subjective rather than quantitative.

Ink Dry Time is a measure of the rate of ink absorption into a sheet of paper. A specific test pattern is printed, the image is blotted, and the
5 resulting optical density of the transferred black ink is measured. The results are fitted to a decaying exponential model from HP, and the ink dry time is calculated. The final dry times are compared to minimum criteria set by HP.

10 Waterfastness is a measure of the amount of colorant transferred from a printed area to an unprinted area when water is applied. The waterfastness test pattern is printed onto the paper, 250 μ l of water is applied across the print, and allowed
15 to run over the printed area and adjacent unprinted area. The optical density of the transferred black ink on the unprinted areas is measured. Resulting optical densities are compared to HP standards.

Highlighter smearfastness is the measure of
20 colorant transferred from a printed area to an unprinted area when contacted by two types of highlighters. The specified test pattern is printed onto the paper and, using an alkaline highlighter and an acid highlighter, a perpendicular line is drawn
25 across the test area. The optical density of transferred black ink on unprinted areas is measured and compared to HP minimum criteria.

A color evaluation is performed with a CIELAB system, which uses an Elrephro Datacolor 2000
30 colorimeter to determine L*, a*, and b* values to score color appearance of cyan, green yellow, blue, magenta, and red printed areas. The results are compared to HP diagrams for color quality.

Toner adhesion is a non-ink jet xerographic
35 test, which measures the degree of toner adhesion to the coating. Toner adhesion is essential for use as a commercial multipurpose paper, which will often be used

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as a copier paper. Here, a test pattern is copied onto the paper using a Xerox copier, the test areas are creased with a 5 pound roller, and the resulting toner loss patterns are compared to a set of standards. The degree of toner loss is rated from 1, best or no toner loss, to 10, worst or severe toner loss.

In the brightness test, the coated paper is tested for TAPPI brightness using the Technidyne S-4 brightness meter. Results are compared to the uncoated base stock.

EXAMPLES

The following non-limiting examples are merely illustrative of the preferred embodiments of the present invention, and are not to be construed as limiting the invention, the scope of which is defined by the appended claims.

Example 1

The preparation of a slurry of calcium hydroxide was carried out by combining 1 part lime with 5 parts water with mechanical agitation at an initial temperature of 50°C. The slurry was mixed for about 10 minutes, diluted with water to 1 part lime to 15 parts water, and screened with a 60-mesh screen to remove grit.

Calcium carbonate was precipitated from the slurry in a 30 liter, stainless steel reactor equipped with variable speed agitation having two pitched blade turbine impellers, heating and cooling jackets, a stainless steel carbonation tube to direct a carbon dioxide containing gas to the impeller, and a probe for monitoring the pH of the suspension.

Twenty-six (26) liters of calcium hydroxide slurry, prepared as described above, were added to the reactor while stirring at 420 r.p.m. To the slurry was then added 6 grams of organophosphonate (Briquest 221-50A, ethanolamine bi-(methylenephosphonic acid), Albright & Williams Americas, Inc., Richmond VA),

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corresponding to 0.1 percent by weight of Briquest based on the calcium carbonate equivalent of the calcium hydroxide slurry. A 1 liter solution containing 95.0 g $\text{Al}_2\text{SO}_4 \cdot 16\text{H}_2\text{O}$ was then added. The
5 slurry temperature was adjusted to 9.5°C., and a gas containing carbon dioxide (28 percent by volume CO_2 in air) was introduced at a rate of 2.3 S.C.F.M. while stirring at 615 r.p.m. Carbonation of the calcium hydroxide slurry was continued for 36 minutes with
10 cooling until a minimum slurry conductivity was reached, indicating that the calcium carbonate precipitation was complete. The final temperature of the slurry was 15.8°C. The specific surface area of the resulting calcium carbonate was determined to be
15 115 m^2/g .

Example 2

In the same manner as described in Example 1, calcium carbonate precipitation by carbonation of a calcium hydroxide slurry with a carbon dioxide
20 containing gas in the presence of 0.1 percent by weight Briquest and 3.15 percent by weight $\text{Al}_2\text{SO}_4 \cdot 16\text{H}_2\text{O}$ was carried out with the same equipment. The carbonation was complete in 35 minutes when a minimum conductivity was attained. The final temperature of the slurry was
25 16.7°C., and the pH was 11.65. The specific surface area of the PCC was 102.3 m^2/g . The PCC slurry was screened with a 325 mesh screen, and placed back into the reactor. While stirring, 45 g of Briquest 221-50A, corresponding to 0.7 percent by weight Briquest, based
30 on the weight of PCC present, was added. The pH was adjusted to 9.5, the slurry was heated to 80°C., and the PCC was heat aged at 80°C. for 5 hours. The specific surface area of the final product was
35 82.4 m^2/g .

Example 3

In the same manner as described in Example 2, a calcium hydroxide slurry was carbonated in the

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presence of 0.1 percent by weight Briquest and 3.15 percent by weight $\text{Al}_2\text{SO}_4 \cdot 16\text{H}_2\text{O}$, where both percentages are based on the calcium carbonate equivalent of the calcium hydroxide. The carbonation time was 39 minutes, the final pH was 10.9, and a PCC with a specific surface area of $115.1 \text{ m}^2/\text{g}$ was obtained. While stirring the mixture, 0.7 percent by weight Briquest 221-50A, based on the weight of PCC was added. The slurry was screened with a 325 mesh screen, returned to the reactor, and heated to 80°C . for 2 hours. The specific surface area of the final product was $81.6 \text{ m}^2/\text{g}$.

Example 4

In the same manner as described in Example 2, a calcium hydroxide slurry was carbonated in the presence of 0.7 percent by weight Briquest 221-50A and 3.15 percent by weight $\text{Al}_2\text{SO}_4 \cdot 16\text{H}_2\text{O}$, both percentages based on the calcium carbonate equivalent of calcium hydroxide. The batch time to minimum conductivity was 36 minutes, and the final pH was 11.4. The specific surface area of the PCC was $99.1 \text{ m}^2/\text{g}$. The slurry was passed through a 325 mesh screen, returned to the reactor, stirred, and 0.15 percent by weight Briquest 221-50A, based on the weight of PCC, was added. The pH of the slurry was adjusted to 9.5, and the slurry was heated to 80°C . The PCC was heat aged at 80°C . for 2 hours, and a PCC with a specific surface area of $63.8 \text{ m}^2/\text{g}$ was obtained.

Example 5

A calcium carbonate slurry with a volume of 0.6 l and a solids concentration of 20.2 percent by weight, based on the total weight of calcium carbonate and water, where the calcium carbonate was a scalenohedral PCC with a specific surface area of $11.57 \text{ m}^2/\text{g}$ and an average particle size of $1.35 \mu\text{m}$ was placed into a 1.5 l stainless steel media mill containing 1.7 kg of glass beads of 0.7 to 0.9 mm. The media mill was

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equipped with variable speed mechanical agitation. The slurry was milled for 10 minutes with an agitation speed of 2450 rpm. The milled product had a specific surface area of 15.59 m²/g and an average particle size of 0.35 microns.

Example 6

A calcium carbonate slurry having a volume of 0.6 l and a solids concentration of 18.7 percent by weight, based on the total weight of the calcium carbonate and the water, where the calcium carbonate was a scalenohedral PCC with a specific surface area of 26.07 m²/g and an average particle size of 0.88 μm, was placed in a 1.5 l stainless steel media mill equipped with variable speed agitation and 1.7 kg of 0.7 to 0.9 mm glass beads. The slurry was milled for 10 minutes using a 2450 rpm agitation speed. The milled PCC product had a specific surface area of 30.11 m²/g and an average particle size of 0.26 μm.

The heat aged and media milled PCC's of Examples 1 to 6 were formulated with an appropriate binder, and paper test sheets were coated with 6 g/m² of the formulation in the manner described above. After drying, the paper hand sheets were tested for true black optical density, composite optical density, spreading, penetration, dry time, and water solubility. The results of these tests are given in Table 1.

TABLE 1

Example No.	1	1	1	2	2	2	2	3	4	5	6
Binder	Starch	PVOH	PVOH	Starch	PVOH	PVOH	PVOH	Starch	Starch	Starch	Starch
Weight Percent Binder	30	15	30	30	15	30	30	30	30	30	30
Compo-site Black Optical Density	0.98	0.9	0.9	1.22	1.11	1.22	1.22	1.15	1.14	1.02	1.03
True Black Optical Density	1.17	1.34	1.31	1.31	1.36	1.54	1.54	1.22	1.25	1.53	1.4
Spreading	106.2	101.3	100.2	108.4	101.6	100.8	100.8	107.7	106.6	101	103.5
Penetration	30.3	34.5	32.7	8.4	25.7	21.8	21.8	20.9	12.1	19.1	23.3
Dry Time	0	0	0	22	0	17	17	11	21	104	50
Water Solubility	0	0.02	0.02	0.15	0.02	0.07	0.07	0.08	0.12	0.45	0.32

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These results demonstrate a significant improvement when compared to a typical uncoated commercial paper, which, typically, has a composite black optical density of about 0.9, a true optical density of about 1.28, spreading of about 108, a percent penetration of about 38, a drying time of about 38, and a water solubility of about 0.17. By comparison, the best high cost, coated commercial paper has a composite black optical density of about 1.2, a true black optical density of about 1.56, spreading of about 98, a percent penetration of about 28, a drying time of about 12, and a water solubility of nearly 0. Therefore, an ink jet recording paper produced according to the present invention, and, in particular, a paper that is coated with a formulation based on the PCC produced in Example 2, is comparable in ink jet print quality to the best high cost commercial paper now available.

While it is apparent that the invention herein disclosed is well calculated to fulfill the objects above stated, it will be appreciated that numerous modifications and embodiments may be devised by those skilled in the art. It is intended that the appended claims cover all such modifications and embodiments as fall within the true spirit and scope of the present invention.

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C L A I M S

1. A process for preparing a precipitated calcium carbonate pigment which comprises forming a slurry of precipitated calcium carbonate particles,
5 adding an organophosphonate compound to the slurry, and heating the slurry to heat age the precipitated calcium carbonate particles therein and form the improved pigment.
2. The process of claim 1, wherein the
10 organophosphonate compound is an amine-containing phosphoric acid derivative and is added in an amount of between 0.1 and 1 percent by weight.
3. The process of claim 1, wherein the
15 organophosphonate compound is ethanolamine bis-(methylenephosphonic acid) and is added in an amount of between 0.4 to 0.85 percent by weight.
4. The process of claim 1, 2 or 3, wherein the precipitated calcium carbonate particles are aged for a
20 sufficient time to impart a BET specific surface area of greater than 60 m²/g.
5. The process of any of claims 1, 2, 3 or 4, wherein the precipitated calcium carbonate particles are aged for between 1 and 10 hours at a temperature of at least 75°C.
- 25 6. The process of any of claims 1, 2, 3, 4 or 5, wherein the precipitated calcium carbonate particles are aged for between about 2 and 5 hours at a temperature of between about 80-85°C.
7. Heat aged calcium carbonate particles
30 whenever obtained by the process of any one of the preceding claims.
8. A process for making an ink jet recording paper comprising: coating at least one side of a paper base stock with a coating comprising heat aged calcium
35 carbonate particles according to claim 7 and a binder to form the ink jet recording paper.
9. The process of claim 8, wherein the coating

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is applied to each side of the paper base stock.

10. The process of claim 8 or 9, wherein the coating is applied to the paper on a papermaking machine during the papermaking process.

5 11. The process of claim 8, 9 or 10, wherein the binder is polyvinyl alcohol or a starch.

12. A process for making an ink jet recording paper having enhanced properties comprising:

a) milling precipitated calcium carbonate
10 pigment to produce particles having a surface morphology and chemistry that enhances the ability of the particles to bind to ink;

b) coating at least one side of a paper base stock with a coating comprising said milled
15 precipitated calcium carbonate pigment particles and a binder to form the ink jet recording paper.

13. The method of claim 12, wherein the particles are milled to have a BET specific surface area of at least $15 \text{ m}^2/\text{g}$.

20 14. An ink jet recording paper comprising a paper base stock, having thereon a coating comprising a heat aged precipitated calcium carbonate pigment, a milled precipitated calcium carbonate pigment, or a mixture thereof, and a binder.

25 15. The ink jet paper of claim 14, wherein the media milled precipitated calcium carbonate pigment has a BET specific surface area of at least about $15 \text{ m}^2/\text{g}$.

16. The ink jet paper of claim 14 or 15, wherein the heat aged precipitated calcium carbonate pigment
30 has a BET specific surface area of at least about $60 \text{ m}^2/\text{g}$.

17. The ink jet recording paper of claim 14, 15 or 16, wherein the binder is polyvinyl alcohol or a starch.

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C09C 1/02, C01F 11/18, B41M 5/00, D21H 19/38	A3	(11) International Publication Number: WO 96/29369 (43) International Publication Date: 26 September 1996 (26.09.96)
(21) International Application Number: PCT/US96/03546 (22) International Filing Date: 14 March 1996 (14.03.96) (30) Priority Data: 08/407,422 17 March 1995 (17.03.95) US (71) Applicant: MINERALS TECHNOLOGIES INC. [US/US]; 405 Lexington Avenue, New York, NY 10174-1901 (US). (72) Inventors: DONIGIAN, Douglas, Ward; 1021 Main Street, Rear, Bethlehem, PA 18018 (US). RESNIK, Robert, Kenneth; 1709 Wynwood Lane, Easton, PA 18040 (US). MCFADDEN, Michael, Gregory; 870 Durham Road, Riegelsville, PA 18077 (US). (74) Agents: RICHARDS, John; Ladas & Parry, 26 West 61st Street, New York, NY 10023 (US) et al.	(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> (88) Date of publication of the international search report: 5 December 1996 (05.12.96)	
(54) Title: INK JET RECORDING PAPER INCORPORATING NOVEL PRECIPITATED CALCIUM CARBONATE PIGMENT		
(57) Abstract An improved thermal ink jet recording paper, incorporating heat aged and/or milled precipitated calcium carbonate ("PCC") and a binder, is disclosed. The heat aging and milling processes produce discrete particles of PCC with a surface morphology and chemistry that enhances the ability of the PCC to bind to ink jet ink without binding so strongly that the color of the ink is changed. Ink jet recording papers incorporating the PCC and binders of the present invention have been produced that are superior to multipurpose office papers with reduced feathering, spreading, and penetration or backside showthrough, as well as improved optical density, dry time, and water fastness. The ink jet recording paper of the present invention is comparable to silica based commercial papers, at a fraction of the cost.		

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 96/03546

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C09C1/02 C01F11/18 B41M5/00 D21H19/38

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C09C C01F B41M D21H

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 344 984 A (PFIZER) 6 December 1989 see claims 1,8 ---	1-7
X	CHEMICAL ABSTRACTS, vol. 116, no. 8, 24 February 1992 Columbus, Ohio, US; abstract no. 62845b, page 197; XP000283529 see abstract & SU 1 650 589 A (VALIULLIN, A. K. ET AL.) 23 May 1991 ---	1,7
A	EP 0 323 697 A (PFIZER) 12 July 1989 see page 4, line 27-29 see page 6, line 40-41; claims 1-3 --- -/--	1-7

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

15 July 1996

Date of mailing of the international search report

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 96/03546

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 055 088 A (PFIZER) 30 June 1982 see page 8, line 14-17; claims 1,2,9 ---	1-7
A	DATABASE WPI Week 9219 Derwent Publications Ltd., London, GB; AN 92-156917 XP002007514 & JP 04 097 910 A (TSUKUMI FINE CERAMI) , 30 March 1992 see abstract ---	1,7
A	DATABASE WPI Week 8541 Derwent Publications Ltd., London, GB; AN 85-252499 XP002007515 & JP 60 166 221 A (MITSUBISHI MONSANTO) , 29 August 1985 see abstract ---	1,7
A	US 5 320 897 A (HIROMASA KONDO ET AL.) 14 June 1994 cited in the application see column 4, paragraph 3 see column 4, last paragraph - column 4, paragraph 1; claims 1,3 ---	8,14,17
A	DATABASE WPI Week 8448 Derwent Publications Ltd., London, GB; AN 84-296922 XP002008161 & JP 59 182 787 A (KANZAKI PAPER MFG.) , 17 October 1984 see abstract -----	8,14,17

INTERNATIONAL SEARCH REPORT

International application No.

PC1/US 96/03546

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

please see attached sheet!

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-11 and 14-17 in so far as linked to 1-11

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/210

1. Claims 1-11 and 14-17 in so far as linked to 1-11.
Process for preparing heat aged precipitated CaCO_3 , thus prepared pigment and process for ink jet recording papermaking
2. Claims 12,13 and 14-17 in so far as linked to 12,13:
Process for ink jet recording papermaking using milled precipitated CaCO_3 and thus obtained paper

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US 96/03546

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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